

SPONTANEOUS ADVERSE EVENTS REPORTING

BACKGROUND OF THE INVENTION

National health authorities and governmental
5 regulatory Agencies in major countries require
Pharmaceutical Manufacturers to collect and report Adverse
Drug Reactions, Adverse Events, Product Complaints and
Device Failure Complaints, and to provide Full Disclosure
about the side effects, risks and contraindications
10 associated with their products. Reports of Adverse Events
are received by pharmaceutical manufacturers via letter,
fax, e-mail or telephone and are usually processed by
safety professionals. The process consists of a series of
sequential and/or concurrent sub-processes: Intake,
15 Investigation and Follow-up, Analysis and Reporting. The
Adverse Event is usually received as a report (Narrative
Verbatim) and is then processed by a healthcare
professional, entered into a database system, catalogued
and linked to a dictionary of international standard
20 medical terminology such as the terms found in MedDRA, the
Medical and Drug Regulatory Affairs Code, or other
nationally or internationally recognized coding schema such
as World Health Organization's Adverse Reactions
Terminology (WHO-ART) Dictionary, or the U.S. Food and Drug

Administration's (FDA) Coding Symbols for Thesaurus of Adverse Reaction Terms (COSTART) schema, either by hand or assisted by a semi-automated processes, called auto-encoding.

5 The investigation and follow-up sub-processes aim at having the event verified and validated by a healthcare professional (physician, hospital). A judgment is reached, whether the Adverse Event report is serious (according to existing definitions established by a national health
10 authority or drug regulatory agency) and unknown, (unlabeled) in which case a report has to be filed with the health authorities within a short period (usually 15 days). All other cases are regularly analyzed and reported on an annual or semi-annual basis in accordance with the laws,
15 regulations or guidelines set forth by a national health authority or drug regulatory agency.

In cases where the Adverse Event occurred in a foreign country or was received by a license taker of the manufacturer, the reports are transferred between
20 manufacturer's safety organizations, requiring rapid information transfer and rigorous quality assurance measures. Special measures have to be taken in cases where reports or data get translated from one language to another. The aim of the reporting process is to discover

previously unknown Adverse Drug Reactions, Drug-Drug Interactions, Drug-Food Interactions or other safety issues at an early stage.

The use of standardized terminology is intended to bring uniformity to the reporting for a large international user base with varied linguistic and cultural differences in reporting Adverse Events and Product Complaints. It is intended as a pragmatic, clinically validated compendium of medical terminology with an emphasis on ease-of-use of data entry, retrieval, analysis and display, with a suitable balance between sensitivity and specificity, within the regulatory environment. A standardized dictionary, such as for example MedDRA is applicable to all phases of development for a drug product and the period during which a drug product is sold, and is also applicable to the product life cycle of most medical devices. By providing one standard of medical terminology, the effectiveness and transparency of medical product regulation worldwide is improved. Standardized terminology for reporting Adverse Events or Product Complaints is grouped in three main associations:

- Equivalence: synonymous terms are grouped together.

- Hierarchical: provides vertical links between super-ordinate (broad grouping terms) and subordinate descriptors
- Associative: allows terms, which are neither equivalent nor hierarchically related but may be related by sign, symptom, body system or organ class, disease and/or diagnosis to be linked horizontally (Special Search Categories).

A standardized dictionary can contain many terms. For example, in the current version of MedDRA at each level is has approximately the following number of terms:

- System Organ Class (SOC) - 26
- High Level Group Term (HLGT) - 334
- High Level Term (HLT) - 1,663
- Preferred Term (PT) - approximately 11,000;
International Level of Communication
- Lowest Level Term (LLT) - approximately 46,000;
Synonyms

The use of multiple standardized terminologies however in reporting and codifying Adverse Events poses challenges and creates coding requirements with respect to accurately

assessing information collected vis-à-vis a single Adverse Drug Reaction or Adverse Event case, or trending information collected across multiple cases albeit for a single drug product or even for similar but not identical drug products that produce the same desired therapeutic effect through similar mechanisms of action within the human body. Because of the large number of terms, there are sometimes too many choices to describe a particular Adverse Event or device complaint. In addition, there are problems due to the fact that each reported event requires a written verbatim narrative of the event, which requires that the safety professional be well versed in the standardized formats and terminologies employed to catalog and characterize various Adverse Events and Product Complaints. Selection of the proper term and creating a highly standardized and structured format applied to each verbatim report are difficult and time-consuming processes requiring great care.

What is desired therefore is a system and method for assisting and rigorously structuring in the intake, investigation and Follow-up, Analysis and Reporting of all Adverse Events and Product Complaints, but especially spontaneously reported cases.

It is also desired to provide a system and method for standardizing Adverse Event and Product Complaint reporting. In that way an untrained health professional or a company employee assigned to receive, process and record Adverse Event reports or can receive and record event reports without the need for intensive training in safety, safety data and information collection, applying and utilizing various coding schema, or applying and utilizing requisite clinical safety knowledge. In addition, with such a system, a patient can, in a highly effective, efficient and standardized manner, report meaningful safety data and information directly to an interested party such as his/her treating physician or a manufacturer of a drug product.

It is also desired to provide a system and method to efficiently generate Adverse Event and Product Complaint reports in order to comply with event reporting time deadlines.

It is also desired to provide a system and method to efficiently, safely and seamlessly transfer the collected data into any standard industry drug safety system, using industry standard protocols and definitions like the E2B and E2C standards defined by the International Conference

on Harmonization of Technical Requirements for Registration
of Pharmaceuticals for Human Use. (ICH M1-3).

It is also desired to provide a system and method to
collect and manage data utilizing a network computer system
operating in a stateless environment, while having the
ability to maintain and manage state.

SUMMARY OF THE INVENTION

The present invention overcomes the drawbacks of the
prior art by providing in a first aspect a method of
dynamically managing and structuring the input and
improving the reliability, accuracy and quality of
collected information by presenting the user with a
hierarchically structured set of predefined terms,
receiving reported information from a user corresponding to
at least one of the predefined terms, categorizing and
characterizing the reported information, dynamically
converting the reported information into a standardized
output and storing the standardized output.

BRIEF DESCRIPTION OF THE DRAWINGS

These and other objects and advantages of the invention will become more apparent from the following detailed description of a preferred embodiment of the invention when taken into conjunction with the drawings

5 wherein:

Figure 1 is an exemplary computer system for implementing the present invention.

Figure 2 is an exemplary computer network system for implementing the present invention.

10 Figure 3 is a flow chart depicting an exemplary site flow diagram for the present invention.

Figure 3a is a flow chart depicting a detailed exemplary site flow diagram for the present invention.

15 Figure 3b is a flow chart depicting a detailed exemplary site flow diagram for the present invention.

Figure 3c is a flow chart depicting a detailed exemplary site flow diagram for the present invention.

Figure 3d is a flow chart depicting a detailed exemplary site flow diagram for the present invention.

20 Figure 3e is a flow chart depicting a detailed exemplary site flow diagram for the present invention.

Figure 3f is a flow chart depicting a detailed exemplary site flow diagram for the present invention.

Figure 3g is a flow chart depicting a detailed exemplary site flow diagram for the present invention.

Figure 3h is a flow chart depicting a detailed exemplary site flow diagram for the present invention.

5 Figure 3i is a flow chart depicting a detailed exemplary site flow diagram for the present invention.

Figure 3j is a flow chart depicting a detailed exemplary site flow diagram for the present invention.

10 Figure 3k is a flow chart depicting a detailed exemplary site flow diagram for the present invention.

Figure 4 is a depiction of an exemplary version of a "Registration" page

Figure 5 is a depiction of an exemplary version of a user login page for the interface of the current invention.

15 Figure 6 depicts an exemplary physician patient consent page.

Figure 7 depicts an exemplary Instructions page for the interface of the present invention.

20 Figure 8 is a depiction of an exemplary depiction of a user identification page.

Figure 9 is a depiction of an exemplary report designation page.

Figure 10a is a depiction of an exemplary Adverse Event reporting page.

Figure 10b is a depiction of an exemplary Adverse Event reporting page.

Figure 10c is a depiction of an exemplary Adverse Event reporting page.

5 Figure 10d is a depiction of an exemplary non-English language Adverse Event reporting page.

Figure 11 is a depiction of an exemplary medication reporting page.

10 Figure 12 is a depiction of an exemplary suspect medication reporting page.

Figure 13 is a depiction of an exemplary medication supplement-reporting page.

Figure 14 is a depiction of an exemplary medical condition-reporting page.

15 Figure 14a is a depiction of an exemplary laboratory test reporting page.

Figure 15 is a depiction of an exemplary written narrative report.

20 Figure 16 is a depiction of an exemplary summary report.

Figure 17 is a depiction of a data flow diagram of the present invention.

Figure 18 is a flow chart depicting the data flow steps of the present invention.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

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I. Introduction

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III. Database Terminology Standardization

IV. Database Encoding

10 V. Embodiments

I. Introduction

The present invention will now be described in detail with reference to the accompanying drawings. While the present invention is described in the context of an Internet based communications network, which includes a specific number and type of components, the system of the present invention may be incorporated into data and voice communications networks of many structures and sizes. The drawings are intended to provide one example of a communication network configuration in which a system of the present invention may be implemented and are not intended to limit the applicability of the present invention to other network configurations. For example, all

of the features of the present invention could be implemented using a voice recognition system, wherein the user could access the system via a landline or wireless mobile communications device and enter information via responses to auditory choices and menus, or to text prompts on a wireless pager.

As will be disclosed below, an aspect of a preferred embodiment involves a process for entering, accessing and encoding data, in any language, through a super-secure portal for Spontaneous Adverse Events Reporting, Medical Product and Device Complaint Reporting Medical Inquiries and registry functions. The Portal is accessed from a Web-Browser via the WWW using strong encryption and HTTPS as the sole protocol between browser and application. No active content is needed in the browser to run the application.

With reference to the various systems and methodologies of the present invention, as described below, aspects of the present invention are described in terms of steps, some of which being executed or executable on a computer system. Various implementations of the inventive systems and methodologies provide a comprehensive, multi-faceted; multi-user based incident reporting system and method. In accordance with these implementations, there is

provided a system and method for entering, accessing and encoding data through a super-secure portal for Spontaneous Adverse Events Reporting, Medical Product and Device Complaint Reporting and Medical Inquiries. In other
5 aspects of the invention, which are discussed in much more detail below, the data encoding routine of the present invention presents a novel method to populate the database in order to provide high accuracy and data entry redundancy.

10 In a preferred implementation, a central database is developed and contains information pertaining to Adverse Events Reporting, Medical Product and Device Complaint Reporting and Medical Inquiries and individual patient information, which preferably is incorporated into the
15 database corresponding to each reported event or complaint. Other types of information can be incorporated, as will become apparent below. Preferably the data fields are named in accordance with the E2B and E2C standards issued by the international committee for harmonization (ICH M1-3).

20 Through the use of the present invention, the process of developing the database incorporates techniques to ensure that the information reported is reliable and accurate. Accordingly, redundant data input systems and methods are incorporated to ensure the reliability and

accuracy of reported information through the various systems and methodologies of the invention. Information concerning the patients' condition, prescribed medication, Adverse Events and complaints are entered into the database through a user interface wherein the user selects information to be included in the database using pull down menus. In addition, once the data is selected, the user must confirm the reported information at least twice. Thus through the immediate review and detection of data input errors, corrective action can be undertaken immediately, by the incident user, therefore increasing the reliability and accuracy of the reported information.

In accordance with one aspect of the invention, patient information, medication information, Adverse Event and Product Complaints are received from the user via the input interface. Data is input through the interface using either pull down menus or free form data entry fields. The pull down menus include a set of clickable data entries for each data categories. In the event that user is unable to locate the data entry from the provided pull down menus that accurately reflects the patients information or condition, then the user can enter that information in a data entry field. However, any data that is entered via the data entry field rather than chosen from a pull down

menu is automatically flagged for verification and validation.

In accordance with one aspect of the invention, patient information is provided and can be accessed by a user or safety monitor through an exclusive password-protected system. In this way, a flexible and paperless reporting environment meets the demands of real-time secure and reliable information needs of patients, treating physicians, clinical investigators and safety monitors within product manufacturers and their licensees and partners. In other aspects, safety reports can be generated from the event reports for use in reporting significant events, or in transferring case reports to another database or data repository.

Computer System Overview

Although a variety of different computer systems can be used with the present invention, an exemplary computer system is shown generally in Figure 1.

Computer system includes a host computer having a processor 204, main memory 208, a secondary memory 210 comprising a hard disk drive 212 and a removable storage

drive 214. The exemplary components of host computer are operably connected via an address/data bus 206, which is not specifically designated. Memory can, and preferably does include a volatile memory (e.g. random access memory), which is coupled with the data bus for storing information and instructions for processor, and a non-volatile memory (e.g. read only memory) coupled with the data bus for storing static information and instructions for processor. Data storage device can comprise a mass storage device.

Host computer constitutes a hardware platform, which executes instructions to implement the application program(s) described below. It will be understood that system, as set forth in Figure 1, is a schematic representation only. Accordingly, the system as described above and below can be implemented as an integral stand alone system as suggested by Figure 1, or can include separate component parts which are interconnected and operable for implementing the invention described below.

Interface device preferably comprises a multi-user network interface (e.g. an Internet interface), which couples computer system to a multi-user system (e.g. the Internet in one embodiment of the present invention). Interface is coupled to permit communication with various application programs contained on the hardware platform

defined by computer system. The Interface can typically be implemented using a graphical user interface (GUI) incorporating pull down menus and data entry fields.

As mentioned above, and in a preferred implementation

5 of the present invention, interface device comprises an Internet interface. The Internet is a well-known connection of worldwide computer systems that operate using a well-known Internet protocol. The Internet is one type of multi-user computer system. Other Internet applications (e.g.

10 using specific protocols) operate on top of the Internet protocol. One such application is the well-known World Wide Web or "www" Internet application that operates using the hypertext transfer protocol or http. The "www" Internet application is a "demand system" in which user requests

15 information from a site and the site transfers the information back to the user on-line. Also well known is the email Internet application which operates using the simple mail transport protocol or SMTP. The email Internet application is a "present system" in that an information

20 transfer command originates from a sender site and information pursuant to that command is presented to the target email address. Another Internet application is the file transfer Internet application that operates using the file transfer protocol ftp. In one embodiment, the present

invention utilizes the www, email, and file transfer
Internet applications as well as the Internet protocol.
Other embodiments of the present invention can be
implemented in other multi-user computer environments. For
5 example, the present invention could be implemented with a
dedicated multi-user system.

Computer system supports a software configuration,
which operates under control of a conventional operating
system. The operating system permits various application
10 processes to be executed. These include, for example, a
communications application that permits data transfer with
various remote terminals as will become apparent below. The
software environment further includes a data management,
storage, and retrieval application that is utilized in
15 connection with data storage device. The data management,
storage, and retrieval application organizes and stores
information, which will be described in greater detail
below. This information is organized and stored within the
environment of the operating system on one or more mass
20 storage devices such as data storage device. Other
applications conventionally known may be included in the
software environment comprising computer system.

In view of the foregoing computer system description
and in accordance with one aspect of the invention, the

reader is referred to Figure 2. There, an exemplary computer system or host system can be seen to comprise part of a network, which includes a database host, a network and a user. In the context of this document, the term "database host" will be understood to include a secure portal hosting the computer program that embodies the current invention, a network server controlled by the server logic (software) and a database for storing site content and input data. The network. Similarly, the term "user" as used in this document will be understood to include a patient, event user, a physician or safety monitor.

The present invention can be implemented with a software program executable on computer system. Such program would typically be stored in memory. A plurality of dictionaries and thesauri are defined in host computer database. Such dictionary and thesauri are stored within a database that is preferably stored within a storage device, such as data storage device. The user interface, as described below includes a plurality of pull down menus.

Each pull down menu contains a list of popular terms used to describe a symptom, medication or indication. The terms contained in the pull down menus are stored in a dictionary and/or thesaurus and are written to the interface display by the software of the present invention. Each of the

popular terms on the user interface is hard coded to a corresponding standardized term. The thesaurus contains information associated with each data entry field of the user interface. Each pull down menu or data entry field that is incorporated into the GUI will be associated with a particular data field in the thesauri. Each data field in the thesauri, corresponding to a user data entry field is also associated with a particular standardized term or code from a standard medical dictionary, such as MedDRA, ICD-9, or WHOART. A data entry contained on the pull down menus is hard coded to at least one field in the thesauri containing a standardized term for that selection. The standardized term, such as from MedDRA, is in turn populated into the database for a particular reported event. In that way it is possible to ensure that the appropriate term is encoded for all entered data. The reported information can include any type of information that is useful in implementing the present invention. Exemplary information includes the user's ID and password, the users reporting status, for example patient, family member, or treating physician, patient information such as age, height and weight, prescribed medication, and categorizations, characterizations and descriptions of an Adverse Event or Product Complaint. Additionally,

particular information regarding the Adverse Event can be included. Exemplary Adverse Event information can include such things as Body Region; Head: Sub Region; Eyes: Symptom; Blurred Vision: Preferred Term; Vision blurred, and various other information as well.

Adverse Event or Product Complaint information is transmitted from a user to a host computer. The Adverse Event or Product Complaint information pertains to at least one event or complaint experienced by the customer. The

Adverse Event or Product Complaint information can be introduced into system in any suitable way. In a preferred embodiment, such information from a user is received electronically, via a suitable data link with host computer, using one or more of the Internet protocols mentioned above. Other methods and systems can, of course, be utilized to permit such information to be received by host computer.

In a preferred implementation, the information that is received into the host computer for each patient pertains to a plurality of different Adverse Events, Product Complaints, medications and symptoms.

The interface of the present invention is described as being presented to the user through the graphical user interface of an Internet browser program. The browser's

graphical user interface (e.g., Netscape™ Navigator™) presents content provided by a resource (e.g., a file) at a URL (Universal Resource Locator). The content can include graphics, text, animation, sound, instructions, etc. A URL
5 can refer to a location on a remote computer that stores the content as data and presentation instruction. The presentation instructions and data can be in a variety of formats such as HTML (Hypertext Markup Language), XML (Extensible Markup Language), PDF (Portable Document
10 Format), JPEG (Joint Photographic Experts Group), and MPEG (Moving Picture Experts Group). When browser requests content from a URL resource, a remote computer providing the resource can transmit the content to a browser for presentation. As shown, the browser is an independent
15 application, however, other applications (e.g., an e-mail program, a word processor, or a spread-sheet) can incorporate functions traditionally performed by the browser.

II. User Interface

20 The Graphical User Interface guides the user through the application through a standardized flow. In the present invention, the user interface is designed so that once the user inputs required information, successive steps are presented automatically. Furthermore, structuring and

standardization of input information are achieved by providing the user with pull down menus for data entry. Free form data entry fields are provided as a backup for the pull down menus. While the interface of the present invention is described as displayed in a web browser window, this description is exemplary of one preferred embodiment of the present invention, those skilled in the art will recognize that many other embodiments with other types of interface elements may be used to provide the same or similar functionality as web browser window.

Figures 3-16 illustrate the data collection and reporting aspects in one embodiment of the present invention. One skilled in the art would appreciate that these various web pages as well as the data fields and links displayed on each page can be omitted, amended, rearranged or adapted in various ways. Furthermore each page may contain cross-referenced links to other pages within the site. In that way the user can directly access a particular page as needed.

Turning now to Figure 3, there is shown a flow chart depicting an exemplary site flow diagram for the present invention. The user upon accessing the site is presented with an index of the site. This includes; the Patient reporting path, the Physician reporting path and the

Monitor reporting path. The index page also contains a link to the registration page for first time users. The user is first presented with a page displaying a site index. The site index page displays information regarding the various reporting section of the interface along with a text, representing a link to each section of the site embodying the present invention. This example Web page was sent from the server system to the client system. This example Web page contains a free form field for entry of users UserID and Password as well as fields to change a users password. In addition there is a link to the registration page for first time users to input registration information. On the Instructions Page, there is included a step-by-step instruction for reporting either; a Side Effect or Adverse Event, or a Product Complaint. The user is directed from the login screen to the patient path, the physician path or the monitor path.

Within the patient path the user is presented with a login screen, a physician patient consent screen, instructions for entering the event report, a screen for the user to indicate his or her identity and a page to initiate either a Adverse Event report or a Product Complaint. To report an Adverse Event the user is presented with a succession of pages to input information

regarding symptoms, medication being taken and herbs and
nutritional supplements being taken. The user is then
requested to identify the medication or supplement that is
suspected of having caused the Adverse Event and the reason
5 that the medication or supplement was taken by the patient.
It would be obvious to one skilled in the art to modify
these text selections choices to be represented as pictures
of medications and supplements. If the user is the
patient, the user is requested to supply information
10 regarding the identity of his or her physician. Likewise,
if the user is a physician, the user is requested to supply
information regarding the identity of the patient
experiencing the Adverse Event.

Upon completion of the data entry, the user is
15 presented with a written narrative and a summary report in
order that the user can confirm the information entered.
Finally the user is asked to confirm his or her identity.

The Product Complaint path is similar to the Adverse
Event path, with the obvious difference being that the user
20 reports information regarding a Product Complaint.

The physician path provides an interface for a
physician to access reports made by or about a patient. By
utilizing this feature of the present invention, a
physician will be able to review reports made by a patient

to correct errors or enter additional information. To use this feature, a physician will, using a password access a list of reports made by or for his or her patients. The list will provide links to the full event reports that the physician can review to either correct or confirm.

Finally, there is provided a monitor path wherein a safety monitor can review all reports for further follow-up as needed. The monitor can view a list of reported cases, which can be indexed and accessed according to well-known principles.

A more detailed view of the interface workflow is shown in Figures 3a-3x. Turning now to Figure 3a, there is shown an exemplary Portal Workflow diagram. The user, upon accessing the site will be presented with a Getting Started page, which has a link to a page presenting patient physician relationship information for the user. After accessing the Getting Started page, the interface business logic directs the user through the registration and login process. After the user has completed the registration and login procedure, the user selects either the Physician, Reporter or Monitor path. As can be seen in Figure 3a, the Physician and Monitor path can alternately follow either a review path, i.e. Physician Confirmation or Monitor, or can be redirected to the Reporter path. The user then proceeds

to enter identity information and indicate if an Adverse Event or Product Complaint will be entered. Turning now to Figure 3b, there is shown a workflow diagram for Adverse Event and Product Complaint reporting path. The interface

5 of the present invention directs the user through the

selected path to collect information regarding the event being reported. Figure 3c depicts the interface workflow for the Physician confirmation path. Upon selecting the Physician Confirmation path, the physician can select a

10 particular patients report from a list and then review and or change data on that report. Additionally, the physician can select, review and verify lab test result information entered by the reporter. Upon completing his review, the physician can confirm the data and submit the data along

15 with an electronic signature. The physician can then review additional patient reports or end his session.

Turning now to Figure 3d, there is shown an exemplary workflow diagram for the Monitor path. Upon accessing the Monitor path, the user can review a list of cases and

20 select from that list. The user can then; search the reported cases for particular data, view and edit previously entered data, input new data and produce case reports.

Turning now to Figure 3e, there is shown a detailed

view of the side effect reporting workflow of the Adverse Event reporting path. The user is directed through the side effect reporting path to provide information regarding the particular body region, sub region, symptom, duration and outcome of at least one side effect.

Turning now to Figure 3f, there is shown a detailed view of the Current Medication reporting workflow of the Adverse Event reporting path. As shown, the user is directed to provided information regarding current

medications, including dosage, formulation, and duration and pharmacy information. Upon completion, the user is can add the medication to a Medicine Cabinet for inclusion in the final report and proceed to Herb and Nutritional reporting or delete the information. In Figure 3g, there is shown a detailed view of the Herb and Nutritional Reporting workflow of the Adverse Event reporting path.

The workflow of this portion of the interface is similar to the Current Medication workflow, however is used to supply information regarding herbs and nutritional supplements

rather than medication. Additionally, upon completion, the user must supply information regarding the medication suspected of causing the adverse event, and the reason for taking the suspect medication. Figure 3h, depicts a detailed view of the Reasons for Medication workflow of the

Adverse Event reporting path. The user is directed to indicate the medication suspected of causing the Adverse Event and the medical condition for which the medication is being taken. Finally as shown in Figure 3i, the user must supply and review additional important information regarding his or her primary care physician or treating physician, reporter information and lab test information.

Turning now to Figure 3j, there is shown a detailed view of the Review Information and Find Out More reporting workflow of the Adverse Event reporting path. As depicted in Figure 3j, the user must review the information supplied, the auto narrative and the summary report. After comparing the report with pre-prepared statistics, the user can enter additional Adverse Event Reports or Product Complaints.

Turning now to Figure 3i, there is shown a detailed view of the Product Complaint Workflow. The user is first directed to provide information regarding the product in question such as medication dosage and formulation.

Additionally, the user is asked to indicate if there is an Adverse Event associated with the Product Complaint. The user can be prompted to enter the Adverse Event data as well as the Product Complaint. If there is no associated Adverse Event, the user is directed to indicate the product

complaint. Product Complaints can be in several categories, for example "Did Not Work"; "Did Not Like", i.e. "Color"; "Something Wrong", i.e. "Package Damaged" and other such information. As in the Adverse Event reporting path, the user must also supply identity information and review and confirm the data supplied.

Turning to Figure 4 there is shown an exemplary version of a "Registration" page. The Registration page displays free form data entry fields for collecting identity information from a first time user. Those fields include in an exemplary version; First Name, Last Name, User ID, Password, Password again, and an additional security devise such as a Secret Question, Secret Answer (a question and answer preselected by the user in order to provide a means to confirm the users identity), phone number and E-mail address. Also included on the registration page will be a statement informing the user that their consent is needed to contact the users physician to gather additional information to complete the report.

The user is required to provide this consent by clicking an Accept button before proceeding with the form. A menu of "Getting Started" steps for a first time or previously registered user is also provided comprising links to the different pages within the site that the user can select to

proceed to the desired login or registration step.

Previously registered users will be directed to a login screen. Figure 5 illustrates the display of a Web page depicting an exemplary version of a user login page for the interface of the current invention. As depicted on the flow chart once the login is complete, the user can select the patient, physician or monitor reporting path.

To begin the reporting process the user is first directed to a page as depicted in Figure 6 which informs the user that both patient and physician identification information is required to adequately complete the reporting process. The user is reminded to include such information on the registration page. The user can then begin reporting an Adverse Event or Product Complaint or complete a report that was previously started by clicking the appropriate button provided.

Upon starting a new event report or restarting a report that was started at a previous time the user is presented with an Instructions page. Figure 7 illustrates the display of an exemplary Instructions page for the interface of the present invention. The instructions indicate to the user the steps needed to report in this embodiment either a Side Effect or Adverse Event, or a Product Complaint, although the instruction page could

include additional information helpful to the user.

Figure 8 is a exemplary depiction of a data input page where the user can select a link or input data by a free form data field or a pull down menu with respect to the his or her identity. The page includes links for Patient Caretaker, Family Member/Spouse, Patient, Pharmaceutical Representative, Treating Physician or Other Healthcare Professional. In addition, there is provided a data entry field for users to enter identity information that is not one of the available categories. Upon selecting one of the links or pull down menu options the user is then directed to the appropriate page for event reporting. In that way the user is directed to the data intake page that is most appropriate to his or her role with respect to the event to be reported.

Turning now to Figure 9, there is depicted an exemplary version of a page wherein the user can designate if the report being made is an Adverse Event or a Product Complaint. After selection of either a Product Complaint or Adverse Event report, additional fields will be presented to the user, wherein additional information such as Age, Height, Weight, Gender and Pregnancy are entered. After entering this data, the user is presented with the appropriate reporting page for either Adverse Events or

Product Complaints.

Figure 10a depicts an exemplary version of an Adverse Events reporting page. On the Adverse Events reporting page the user has several fields in which to input Adverse Event information. Included in the data that the user must input are the Adverse Event symptoms. The symptoms list contains popular names for events that a patient would use to describe their condition. The user can scroll through the list to select the symptom or symptoms that most accurately depict the Adverse Event. In addition, there is provided a diagram depicting a human body. The diagram is divided into regions that represent links for displaying symptoms for each region of the body. For example, one can select the shoulder and arm region link on the body diagram. After selecting a particular region, a sub region list will be presented to the user, as shown on Figure 10a, where the user can more specifically delineate the region of concern. For example the user can designate the right, left or both sides of a selected region. Upon designating a body sub region there will then be displayed symptoms, which are pertinent to that body sub region that the user can select by clicking as shown on Figure 10b. The symptoms are listed using popular names for each symptom. Each popular symptom description is linked or hard coded to

the standardized medical dictionary term and code for that symptom. As shown on Figure 10c, the user is asked to indicate the duration of any symptoms by inputting the date that any symptom started, the date that a symptom ended and further indicate if the symptom is ongoing. The user must also enter information regarding the end result of the Adverse Event such as hospitalization or death of the patient. Finally, the user is asked to provide information regarding what action was taken in response to the Adverse Event. The user is presented with several choices that can be checked to indicate a particular action, such as Did Nothing, Consulted a Physician, Stopped Medication, Reduced Dose or Switched Medication. With respect to the Reduced Dose or Switched Medication data input choices, the interface provides the user with free form data entry fields for providing the necessary information. Furthermore the user is provided with a check box to affirm if the actions helped, a check box to indicate if the effect returned when medication was resumed, if another medication was taken against the Adverse Effect and a check box to indicate if the additional medication was effective. Finally there is provided a free form data entry field to indicate any alternate remedies that the patient pursued. Of course this data entry selections and graphical diagrams

can be modified to customize the interface for use in a particular application. One such modification is depicted in Figure 10d. Figure 10d shows an exemplary adverse events reporting page displayed using the German language.

5 The interface of the current invention can be modified to display all text in any language. Furthermore, the interface can be modified to display pharmaceuticals and products available in a particular market. In addition to changing the language, the interface can be further
10 modified to reflect local differences in pharmaceutical products, description of symptoms, and other differences. Therefore, Figure 10d is meant to exemplary only, one skilled in the art could see that it would be possible to incorporate a change in the language of the interface, as
15 well as other modifications while maintaining the functionality of the invention.

Turning now to Figure 11, there is depicted a data entry page for the user to enter information regarding the medications that the patient is taking. There is provided
20 an index of medication names that the user can click on in order to select the medication that they are taking. Alternately the user can select the medication from a pull down menu or enter the medication name in a free form data entry field. The medication list is comprised of all

commercially available medications. In addition, each listed medication can include a graphic depiction of the medication, in order to assist the user in selecting the correct medication from the pull down menu. The user must also provide the dosage and formulation information by selecting from a pull down menu.

After the user has selected the medication, dosage and formulation information and provided the information regarding length of usage time and optional information the selection is recorded and displayed in a data field on the page known as Your Medicine Cabinet by the user clicking on a button to add the medication to the list of medications. Once the user has entered a first medication into the Medicine Cabinet, he or she can repeat the process to add another medicine, delete a medicine from the list by selecting and clicking on the delete button or indicate that he is finished entering medications to the list by selecting the Done button. Additionally in Figure 12 there is shown an exemplary web page where a user must select the medication or medications, which are suspected of causing an Adverse Event. Displayed on the page is a list of the medications that the user has previously indicated are being taken. Each medication listed is listed together with a check box. The user is prompted to enter a check in

the box of for each medication that is believed to have caused an Adverse Event. Once the user has indicated which medication is suspected of having caused an Adverse Event, additional information is requested from the user regarding the medical condition of the patient with respect to the use of the medication. In that way it is possible to include in the database information regarding what is known as "off-label" use of a medication, i.e. use of a medication for a non-indicated condition.

Similarly as depicted in Figure 13 there is provided a data entry page having the same data fields and as that depicted in Figure 11, however it is intended for the user to provide data related to herbs or nutritional supplements that the patient may also be taking in addition to any prescribed medication. This information is important in evaluating the Adverse Event, as there may be a contraindication between a medication and a nutritional event.

Turning to Figure 14 there is shown a web page where a user can indicate the medical condition for which a medication has been prescribed. The medications that the patient is taking may have been previously indicated as the suspected source of the Adverse Event are listed on the page; the user can then select from a pull down menu the

medical condition for which the medication is prescribed.
The pull down menu lists only the on-label uses for each
medication. If the condition is not listed on the pull
down menu, then the user can enter the condition via a free
5 form data entry field. As described hereinafter, when a
condition is entered in the data entry field, the entry is
flagged in the database for further review by a safety
monitor. After the user enters the information indicating
the disease for which the medication is prescribed, they
10 are then directed to either the patient or physician
information page. If the user is the patient, then the
patient information would have already been entered and the
user will be directed to the physician information page,
conversely if the user is the physician, then the physician
15 information would have already been entered and the user
will be directed to the patient information page.

In the same way that the information is gather
regarding Adverse Event symptoms and medication, the
interface of the present invention could be modified to
20 further include web pages to collect information regarding
any diagnostic tests performed on the patient with respect
to the Adverse Event. This could include laboratory tests.
Figure 14A depicts an exemplary page for reporting the
results of Laboratory test results associated with the

report of an adverse event or product complaint. Of course the method of the present invention is not limited to collecting information regarding any particular diagnostic test and could be modified to accommodate a particular
5 circumstance.

Turning now to Figure 15, there is depicted a page displaying a verbatim narrative that is compiled from the information provided by the user. The narrative is generated by the narrative engine that is part of the
10 software embodying the present invention. The narrative engine uses the response that the user has input and formulates a highly structured and standardized narrative of the Adverse Event. The narrative engine employs programming logic that would be well known to one skilled
15 in the art. The narrative provides an additional means of quality control for the event report by reiterating the users input in a standardized, structured, clear and concise manner. The user is given an opportunity to review all of the information provided in the form of a written
20 narrative and add or correct any information.

After reviewing the narrative report, the user is given a final opportunity to verify the information provided. Figure 16 depicts a page displaying a summary report of all information previously entered. The

information includes Patient Information, Adverse Event information, Suspected Medications and Concomitant Medication. The user is instruction to type over text to edit any information appearing in the summary and then
5 indicate that the report is complete by clicking on a button provided. Finally the user is presented with a page displaying data fields for entering identification information for the user. Information such as user name, address, telephone number, date of birth and occupation are
10 typically included.

In accordance with the description of the above information gathering pages of the interface of the present invention, there is also provided within the patient reporting process a path to report Product Complaints.

15 While the information gathered will not be exactly the same as that collected for an Adverse Event, modifications to the interface and database of the present invention will be obvious to one skilled in the art. Likewise the interface also provides a physician path and a monitor path. These
20 paths differ from the patient path in that the main function of each is to provide information to a safety monitor or a patient's physician. Therefore the interface is adapted to report information that has been previously entered in the physician and monitor path.

III. Database Terminology Standardization

The Adverse Event and Product Complaints that are furnished by the user are input using common popular terms for each symptom in accordance with the vernacular language of a particular country or culture. It is however, necessary to describe all symptoms in standardized terms in the final report the pharmaceutical company uses in assessing an event. For example, symptoms can be described using MedDRA terminology to ensure that like events are described in like manner. However, due to the large number of terms in standardized dictionaries such as MedDRA, it is difficult to efficiently encode the description of an Adverse Event or Product Complaint.

The standardized terms that are presented to the user are compiled in such a way so as to correspond to everyday common terminology used to describe symptoms. Thus rather than utilizing a symptom list derived from a standardized industry dictionary such as MedDRA, the symptom list is compiled by first generating a list of commonly used terms to describe a symptom. Those terms are then hard coded to correspond to a single particular standardized symptom term. In this way, the user of the present invention is presented with an interface that is easier to use and thus

results in more reliable reporting by the user.

IV. Database

Turning now to the database component of the present invention, there is provided a database with a software component for managing and storing data that is transmitted from a user to the host computer embodying the present invention. Utilizing the Internet to collect and manage data allows an enterprise to quickly and efficiently connect to many and distant remote users, however, a number of problems occur when using a Internet base data management system which are not addressed by the current systems.

A data management system utilizing Internet-based communication is different from mainframe and client-server arenas traditionally used for data management. One difference is that HTTP, an underlying protocol of internet communication, is both connectionless and stateless.

The Internet is said to be stateless because no information about what occurred previously is maintained. Most modern applications maintain state, which means that they remember what you were doing last time you ran the application, and they remember all your configuration

settings. This is extremely useful because it means you can mold the application to your working habits.

The World Wide Web, on the other hand, is intrinsically stateless because each request for a new Web page is processed without any knowledge of previous pages requested. This is one of the chief drawbacks to the Internet and the HTTP protocol. Because maintaining state is extremely useful, programmers have developed a number of techniques to add state to the World Wide Web. These include server APIs, such as NSAPI and ISAPI, and the use of cookies.

The stateless condition of the Internet causes a problem for dynamic interactions where a web system needs to be able to keep track of the user's state during a session involving multiple web interactions (e.g., web page requests). Without a way to manage state, between web transactions the system will have "forgotten" information about the user and the context of the session. This can be further complicated by the fact that in many large web systems the user does not interact with the same web server during a session.

The stateless condition is especially a problem when using the Internet for collecting data. When collecting

complex data using an Internet based application, it is usually necessary to have multiple web pages within a site for identifying and registering a user, collecting data and confirming user input. This is a problem in a stateless environment, because information entered identifying the user or data collected on one page is lost when the user directs his browser to another page, unless that information is saved directly to a database or cookies are used. Saving information directly to a database as soon as it is entered is an inefficient process and can result in incorrect data being included in the database.

The present invention overcomes the traditional problems of using a stateless environment for data management by utilizing a software data management tool for receiving, managing and storing data.

All data collected from the user in the current invention is stored in a database that includes the user and patient identity information as well as the incident report. The database makes use of an XML encoding because of its capability to create large repositories of data that operate across any computer platform. For example one can map the fields of a database to elements, and then share that information across a vast array of platforms. The XML database structure is hosted on the server side and then

the XML is completely decomposed into HTML interface components (form elements) at the application server and sent as HTML to the client where it is rendered.

The XML data structure provides an extremely flexible
5 architecture for structured information collection,
manipulation and storage. The interface portal of the
present invention implements a data collection mechanism,
which moves users from page to page, enforcing business
logic (such as ensuring that required fields have content)
10 and populating that data to an XML structure. As each page
is completed, the XML structure is serialized and moved to
the next page where it is deserialized and the next page's
content is loaded into the structure. When a person has
completed the required information, the XML structure is
15 deserialized, each data field is inspected to ensure that
the data conforms to specified constraints, and the
information is entered into the system. The entire
populated XML structure is also stored in the database and
used to quickly populate screens when monitors or other
20 medical personnel request the file.

The flexibility provided by using XML to manage the
data allows the Portal and collected data to be highly
modifiable. While data is deserialized and stored
discreetly in the database, that data is really only used

for quick searches. The actual work is performed through the passing, storage, and manipulation of the XML structure.

Furthermore, in another embodiment, it would be possible to implement the present invention by utilizing signing certificates. It would be possible to actually send a user the XML structure so they can sign it. The signed objects could then be stored in the back end for non-repudiation.

Turning to Figure 17, there is depicted a block diagram representing components of the system of the current invention for the transfer and storage of data. The system is depicting as including a user interface, a software package, commonly referred to as "middleware" that controls the flow of data between the user and the host, including logic and data checking. In addition there is provided a thesauri database, a user database, and an event report database.

There is shown in Figure 17 an exemplary interface screen, the interface software and logic, and a plurality of database storage media. The interface includes a plurality of data entry categories such as symptom, medication-dosage, and medication-indication. A user enters data using the interface through a plurality of pull down

menus or free form data entry fields. It is preferable that the user enter all data by selecting an entry from the provided pull down menus, therefore it is a feature of the present invention to include a full range of pull down
5 menus to cover a majority of the possible data entries for each category. In that way it is possible to reduce input errors by providing the user with a list of the most common data entries. Therefore a user will only need to enter data without the aid of a pull down menu in exceptional
10 circumstances.

The interface software displays the information to the user through the interface of the present invention in the exemplary version described herein. A web page text file incorporating a set of HTML tags describes how the text
15 should be formatted when a browser displays it on the screen. The tags are simple instructions that tell the Web browser how the page should look when it is displayed. A Web browser, like Netscape Navigator™ or Microsoft Internet Explorer™, requests a page, so that the browser can pull
20 the page through the network and into the users machine and interprets the set of HTML tags within the page in order to display the page on the users screen. Additionally, the interface software includes logic, which controls the flow of data to and from the user interface. The interface

logic includes a dynamic XML component that reads data entry choices from the database, displays those choices on the interface and records the users selections on a dynamic web page. Once the user has confirmed the selections the

5 dynamic XML component parses the data to a storage database. The process can be described with respect to the Figure 18. In step 1, the user accesses a web page of the current invention, wherein the dynamic XML component refers to the thesauri database. The XML component reads the

10 appropriate common language term to be displayed in each pull down menu field of the accessed web page. Each common language term is linked to the appropriate medical dictionary term and code in a corresponding thesaurus of the database. In step 2 a user selects a data entry term

15 from the plurality of data terms listed in a particular pull down menu. The dynamic XML component, in response to the selection made by the user, reads the user selection, from the interface and writes that data into a dynamic XML page together with the corresponding term and code from the

20 linked thesauri. The dynamic web page is established by the dynamic XML component for each event report initiated by a user, and is provided to temporarily maintain the data until the user has completed all data entry. In that way any changes or edits can be easily made prior to writing

the data to the storage medium of the database. By
utilizing the dynamic XML component all logic can be
contained in the dynamic XML component, and the database
can be a simple storage device. Therefore, any changes
5 with respect to how the data is saved can be implemented in
the XML component without the need to change or amend the
database. In alternate step 2 the user may provide a
response to an information request by manually entering the
data in a free form data field. In the exemplary version
10 described here as a web based user interface, the user
would position the cursor into a free form data field and
type the data entry. The XML component then reads the data
entered and writes the response to the dynamic XML page.
The XML component also flags certain response for further
15 review by a safety monitor. Those responses include;
manually entered responses, and responses which indicate
that the reported event may be potentially serious and/or
non-labeled. The seriousness of an event is determined by
automatically assessment of the selected outcomes vs. a
20 standard list of "serious" outcomes as defined by the
regulatory bodies. The "labeled-ness" of the event is
automatically determined by comparing the selected symptoms
with a pre-defined list of MedDRA or other schema of
preferred terms, characterized by the manufacturer as

"labeled" or "potentially labeled". In that way an additional quality control check is added for data that is manually entered by the user that may be prone to errors, inconsistencies and/or inaccuracies. Steps 2 and alternate
5 step 2 are repeated by the user until all data has been entered. The XML component will write each data entry to the dynamic XML page. In step 3 the user is presented with a web page that displays all data entered with respect to the event being reported. The user can then confirm or
10 edit the displayed responses and then enter a final confirmation by clicking a provided button. After the user issues a final confirmation, the XML component will then parse the data, step 4, held on the XML page into the appropriate data fields in the database where the data is
15 permanently stored.

The data is permanently stored using a database server. The database server is an application program that interacts with a database. Generally, the functions of the database server are to receive requests for information
20 directed to the database, format the requests into queries that are understood by the database, submit the requests to the database, receive one or more results from the database, format the results, and deliver the formatted results to one or more other application programs. For

example, the database server may receive a request for information in the database in the form of a Structured Query Language (SQL) statement that identifies one or more tables in the database. The database server verifies that the SQL statement has correct syntax and identifies information that is actually in the database. The database server submits the SQL statement to the database and receives a result set of records from one or more database tables.

10 The browsing engine is an application program that interacts with the other application programs to carry out a method of browsing a hypertext system in the manner described further herein. Generally, the browsing engine establishes one or more connections with one or more clients, presents a view of hypertext documents in the database to the clients, receives information that identifies which documents in the database are relevant to the clients, creates a ranked list of the documents based upon the relevance information, and provides the ranked list to the clients. Other functions of the browsing engine are apparent from the description herein.

V. Embodiments

In an alternate embodiment, the present invention can

be utilized to automate the process of collecting data having significance to a particular group of users. For example, the present invention could be used to report problems with consumer goods, such as automobiles.

5 Complaints and problems with new automobiles are monitored by the manufacturer for quality control, by consumer groups to gauge value, and by the federal government for recall purposes. In an alternate embodiment of the present invention, certain aspects would be adapted to the
10 alternate use. For example, the interface and database of the current invention could be adapted for use in receiving reports on automobile defects and problems.

With respect to the user interface, the user would be presented with pages containing images and information
15 consistent with the purpose of the site. For example the network site embodying the present invention could include a consumer path a manufacturer path and a monitor path. The site could provide a means for a consumer to easily and efficiently report problems regarding a vehicle to the
20 manufacturer. In this alternate embodiment, a consumer path would replace the patient path. Figure 10a,b,c would alternately depict a diagram of a generic vehicle, which the user would be able to click on to select various systems and sub-systems. As in the current embodiment,

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10 selecting a system or sub-system, would display to the user
a plurality of pull down menus containing a list of
problems associated with that area of the vehicle that the
user could click on in order to report a problem or
5 difficulty. It is obvious to one skilled in the art that
the system embodying the present invention could be further
modified to accommodate this or other alternate
embodiments. Other alternate embodiments could include a
system for reporting problems with consumer goods other
10 that motor vehicles and health or disease control problems
with respect to animals. Other uses and modifications to
the present invention will be obvious to those skilled in
the art.

The techniques described here are not limited to any
15 particular hardware or software configuration; they may
find applicability in any computing or processing
environment. For example, functions described as being
performed by a server can be distributed across different
platforms. Moreover, the techniques may be implemented in
20 hardware or software, or a combination of the two.

Preferably, the techniques are implemented in computer
programs executing on programmable computers that each
include a processor, a storage medium readable by the
processor (including volatile and non-volatile memory

and/or storage elements), at least one input device and one or more output devices. Program code is applied to data entered using the input device to perform the functions described and to generate output information. The output
5 information is applied to one or more output devices.

Each program is preferably implemented in a high level procedural or object oriented programming language to communicate with a computer system, however, the programs can be implemented in assembly or machine language, if
10 desired. In any case, the language may be a compiled or interpreted language.

Each such computer program is preferably stored on a storage medium or device (e.g., CD-ROM, hard disk or magnetic diskette) that is readable by a general or special
15 purpose programmable computer for configuring and operating the computer when the storage medium or device is read by the computer to perform the procedures described in this document. The system may also be considered to be implemented as a computer-readable storage medium,
20 configured with a computer program, where the storage medium so configured causes a computer to operate in a specific and predefined manner.

Thus, the foregoing descriptions of specific embodiments of the present invention have been presented

for purposes of illustration and description. They are not intended to be exhaustive or to limit the invention to the precise forms disclosed, and obviously many modifications and variations are possible in light of the above teaching.

- 5 The embodiments were chosen and described in order to best explain the principles of the invention and its practical application, to thereby enable others skilled in the art to best utilize the invention and various embodiments with various modifications as are suited to the particular use
- 10 contemplated. It is intended that the scope of the invention be defined by the Claims appended hereto and their equivalents.